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**APPLICATION
FOR
UNITED STATES LETTERS PATENT**

Title: ACTIVIATION OF ELECTROCHEMICAL
CELLS WITH CATALYST ELECTRODES

Inventors: Zhigang Qi
Arthur Kaufman

ACTIVATION OF ELECTROCHEMICAL CELLS

WITH CATALYST ELECTRODES

Field of the Invention:

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This invention relates to electrochemical cells, such as proton exchange membrane (PEM) fuel cells, direct methanol fuel cells (DMFC) and, more particularly, to electrochemical cells operating at ambient temperatures and pressures and which comprise membrane-electrode assemblies whose electrodes are lightly loaded with catalyst.

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BACKGROUND OF THE INVENTION

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The performance of a proton exchange membrane (PEM) fuel cell and a direct methanol fuel cell (DMFC) is largely determined by the membrane-electrode assembly (MEA), which is composed of an anode that oxidizes the fuel, a cathode that reduces oxygen, and a proton transferring, ion-conducting membrane. The membrane prevents electrical short-circuiting between the anode and the cathode, and separates the fuel from the oxidant.

A PEM fuel cell normally uses hydrogen as the fuel, while a DMFC uses methanol as the fuel. The terms "fuel cell", or "cell", as used hereinafter, shall define either a PEM or DMFC-type fuel cell.

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The fuel oxidation reaction and the oxygen reduction reaction of these cells, typically, are kinetically slow. Therefore, catalysts such as platinum and its alloys are often used to speed up these reactions.

PLATINUM ALLOY CATALYST

10 In the fuel cells, the catalysts are generally made into porous layers that serve to increase the contact area between the reactants and the catalyst particles. The layers can be applied directly to the membrane or they can be applied to a 15 gas diffusion medium. Carbon paper and carbon cloth-type materials are commonly used as gas diffusion media because of their good electrical conductivity, high corrosion resistance, and controllable porosity.

20 It is desirable to decrease the amount of noble metals used to fabricate the fuel cells in order to reduce the cost of manufacture. Catalyst layers comprising metal black with lower surface areas were initially used in the fuel cells.

The catalyst loading for an electrode was required to be over 4.0 mg/cm² in order to achieve good performance.

Subsequently, catalyst layers containing supported metal nano-particles featuring higher surface areas were prepared, as

5 illustrated in United States Patent Nos. 4,166,143, granted to

Petrow et al on August 28, 1979, 4,876,115, granted to

Raistrick on October 24, 1989, and Re. 33,149 granted to

Petrow et al on January 16, 1990. The support has several

functions. It provides sites for anchoring the metal

particles during and after their formation. The particles

formed in such an environment have small and uniform size.

Particle coalescence, or aggregation, becomes less likely.

Therefore, the catalyst does not lose its surface area as fast as an unsupported catalyst due to the chemical/physical

15 interaction between the support and the metal particles. The

support also provides electrical connection to catalyst

particles carried upon different support materials. Carbon

black is the most practical support material used in fuel

cells because of its large surface area, good electrical

20 conductivity, and high corrosion resistance.

Electrons and protons are both involved in fuel cell reactions, so it is necessary to provide good electrical and

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proton conductivities. This limits the reaction zone within the interface between the catalyst layer and the ion-conducting membrane for a traditional MEA. This interfacial region is extremely thin and the total surface area of the catalyst particles in this region is low. Thus, the catalyst layer cannot provide a high current density. The catalyst that is not in contact with the membrane is simply wasted.

This can be changed by incorporating an ionic conductor, such as Nafion®, a perfluorinated ionomer made by E. I. DuPont, into the catalyst layers.

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After Nafion incorporation, the entire catalyst layer conducts both electrons and protons so that catalyst utilization in the layer is improved dramatically. The catalyst layer will in turn generate and sustain a higher current density. Nafion can be impregnated into a catalyst layer by brushing and spraying it upon an electrode surface or by respectively floating or dipping the electrode upon or into a Nafion solution. Some illustrations were made by Ticianelli et al., Methods to advance technology of proton exchange membrane fuel cells, J. Electrochem., Soc. pp. 2209-2214 (1988), September; and Poltarzewski et al., Nafion® distribution in gas diffusion electrodes for solid polymer-

electrolyte-fuel-cell applications, J. Electrochem. Soc. pp
761-765 (1992), March.

The advantage of applying Nafion this way provides an
5 opportunity to incorporate polytetrafluoroethylene (PTFE),
which is a water repelling agent, into the catalyst layer.
This is usually accomplished before the application of Nafion
so that the final catalyst layer will have a controllable
10 hydrophobic character in order to reduce the likelihood of
flooding. The disadvantage of this method, however, is that
it is very difficult to control the amount of Nafion that is
applied. This results in a non-homogeneous distribution of
the Nafion material over the entire catalyst layer. The
regions containing more Nafion material will be easily
15 flooded, while the regions with insufficient Nafion material
may not be able to provide enough proton conductivity. The
Nafion may localize on the surface in one place, but penetrate
the underlying gas diffusion medium in another place.

20 Another method of incorporating Nafion into the catalyst
layer is to mix the catalysts directly with the Nafion,
especially supported catalysts, and then use the resulting
mixture to fabricate the catalyst layer, as illustrated in

United States Patent Nos. 5,211,984, granted to Wilson on May 18, 1993, 5,723,173, granted to Fukuoka et al on March 3, 1998, 5,728,485, granted to Watanabe et al on March 17, 1998, and 6,309,772, granted to Zuber et al on October 30, 2001.

5 The catalysts and Nafion mixture is easily achieved; they can
form an even distribution through the entire catalyst layer.

Solvents such as glycerol may be used during the mixing in

order to achieve good viscosity and hold the catalyst

particles in suspension in order to minimize their

agglomeration, as shown in U.S. Patent No. 5,211,984, granted to Wilson on May 18, 1993.

Sometimes, the Nafion solution is converted into a colloidal suspension by adding a proper organic solvent before mixing it with catalysts, as illustrated in United States Patent No. 5,723,173, granted to Fukuoka et al on March 3, 1998. This patent teaches that such a colloidal suspension can provide a good Nafion network for achieving a uniform distribution with the catalyst particles.

Directly mixing the Nafion with the catalyst, however, makes it almost impossible to incorporate PTFE into the catalyst layer. This is because PTFE needs to be sintered at

a temperature in excess of 330°C, but such an elevated temperature will destroy Nafion. Without the PTFE, the catalyst layer is more likely to be flooded.

5 All of these recent developments have helped to decrease the catalyst loading from 4.0 mg/cm² or higher to 0.5 mg/cm² or less. However, a fuel cell that has slightly loaded catalyst electrodes and operates under ambient conditions shows much lower performance than a fuel cell using highly loaded catalyst electrodes. Even though the catalyst layer is made to conduct both protons and electrons through the mixing of the supported catalyst with Nafion, many of the catalyst sites will not become active under ambient conditions.

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15 The present invention provides a procedure and an article made by the procedure, whereby catalyst utilization can be increased. The method comprises the activation of the membrane-electrode assembly. The activation procedure is much shorter than traditional "break-in" processes, yet also dramatically increases catalyst utilization.

SUMMARY OF THE INVENTION

Mixing a proton conducting material, such as Nafion, with a catalyst layer will improve fuel cell performance. However, 5 providing a three-dimensional catalyst layer to conduct protons will not necessarily provide a well performing electrode. Many of the catalyst sites in such a layer will not be available for reaction. Some of a number of reasons could be that: a) the reactant cannot reach the catalyst sites because they are blocked; b) the Nafion near these catalyst sites cannot be easily hydrated; or c) the ionic or electronic continuity is not established at these sites.

Catalyst sites that cannot participate in the 10 electrochemical reaction of the fuel cell are "dead". Making 15 these "dead" sites active is one of the objectives of this invention. It has been found that temperature and pressurization play a crucial role in converting "dead" sites into active sites. Operating a fuel cell above ambient 20 temperature, preferably close to 100°C, and applying a few atmospheres of pressure to the gaseous reactants, will quickly provide an activated fuel cell after only a number of hours. Such an activation procedure results in a dramatic increase in

catalyst utilization. The fuel cell can then generate a much higher current density under ambient conditions.

One objective of the present invention is to provide
5 increased catalyst utilization by a new activation procedure,
especially for the electrodes of fuel cells with low catalyst loadings and fabricated with supported catalysts.

Another objective of this invention is to provide a means
10 to boost the MEA of a fuel cell to a high performance within a few hours.

Still another objective of the present invention is to provide a fuel cell starting procedure, especially one where
15 the fuel cell is intended to be operated under ambient conditions (e.g., ambient temperature and pressure).

BRIEF DESCRIPTION OF THE DRAWINGS

A complete understanding of the present invention may be obtained by reference to the accompanying drawings, when

considered in conjunction with the subsequent detailed description, in which:

5 FIGURE 1 shows a graph of the performances of an electrode (cathode) with a Pt loading of 0.12 mg/cm² at a cell temperature of 35°C before and after the activation procedure of this invention;

10 FIGURE 2 illustrates a graph of the performance of the activated electrode of FIGURE 1 at different temperatures;

15 FIGURE 3 depicts a graph of the performances of the activated electrode shown in FIGURE 1 within a four-week period after the initial activation;

FIGURE 4 shows a graph of the performances of an electrode after being activated for different lengths of time;

20 FIGURE 5 illustrates a graph of the performances of various Pt-loaded electrodes before activation; and

FIGURE 6 depicts a graph showing the performances of the electrodes shown in FIGURE 5 after activation.

PREFERRED EMBODIMENT OF THE INVENTION

Generally speaking, a fuel cell is shown that is operated at a temperature and pressure for converting "dead" sites in
5 the catalyst layer into active sites. Operating a fuel cell above ambient temperature, preferably close to 100°C, and applying a few atmospheres of pressure to the gaseous reactants, will quickly provide an activated fuel cell after only a number of hours. Such an activation procedure results in a dramatic increase in catalyst utilization. The fuel cell can then generate a much higher current density under ambient conditions.

EXAMPLE:

An electrode with platinum loading of 0.12 mg/cm² was made using E-TEK 20% Pt/Vulcan XC-72 material. The catalyst and Nafion solution were mixed thoroughly before being applied to a gas diffusion medium, ELAT, a carbon cloth impregnated with carbon black and polytetrafluoroethylene (PTFE). Two pieces of the electrode with an active area of 10 cm² were hot-pressed onto a Nafion 112 membrane at 130°C.

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The resulting membrane-electrode assembly (MEA) was tested in a single cell test fixture. The test fixture was composed of a pair of metal plates with serpentine flow-fields. The plates were coated with a metal nitride for corrosion protection. Rod-like heaters were inserted into the plates to control the cell temperature. Air and pure hydrogen were used as the reactants. The gaseous reactant, when needing humidification, was passed through a stainless steel water bottle. Otherwise, the gas bypassed the humidifier and was introduced to the cell at room temperature. A cell temperature of 35°C, a hydrogen humidification temperature of 45°C, and air humidification temperature of 45°C, are denoted herein as 35/45/45°C.

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Now referring to Curve A of FIGURE 1, the performance of the electrode is illustrated after the cell has been operated at 35/45/45°C for 180 minutes. The performance increased gradually during this time period, but after 180 minutes the cell did not show further increase in performance. The cell seems to reach its highest performance, no matter how much longer it is run under these conditions.

Curve B shows the performance of the same electrode after activation for 120 minutes. The activation was performed at a cell temperature of 75°C, hydrogen humidification temperature of 95°C, an air humidification temperature of 90°C, a hydrogen backpressure of 20 psig, and an air backpressure of 30 psig.

The activation condition was denoted at 75/95/90°C and 20/30 psig. During the activation, the cell voltage was set at between 0.40 and 0.60 V, for most of the time, in order to sustain a current density of 1.0 to 1.5 A/cm². However, the load was periodically changed so that the cell voltage was changed from open circuit voltage to as low as 0.1 V, backward and forward a few times. The cell performance at 75/95/90°C and 20/30 psig was recorded every 30 minutes until it did not show any further increase. The temperatures of the cell and the humidifiers were then decreased to 35 and 45°C, respectively. During this cooling process, no load needs to be applied. After the cell and the humidifiers reached the desired temperatures, a load was applied and the performance of the cell at 35/45/45°C was recorded. This was the performance of the cell after the activation. A dramatic increase in performance was achieved after the activation. Such an increase could never be achieved if the cell were

tested only at 35/45/45°C without the activation procedure of
this invention.

Referring to FIGURE 2, the performance of the activated
5 electrode of the Example is shown at different cell and
humidification temperatures. The conditions for Curves C, D,
E, and F were 35/45/45, 50/65/60, 60/80/75 and 75/95/90°C,
respectively. The humidification temperature was increased as
the cell temperature was increased in order to make sure that
the MEA was fully hydrated. Although the reactants'
concentration, or partial pressure, decreased with increase in
humidification temperature due to the presence of more water
vapor, the cell performance increased with the cell
temperature. Evidently, the increase in the kinetics of both
15 the oxygen reduction reaction and hydrogen oxidation reaction
with temperature overcame the decrease in the reactants'
concentrations.

The activation effect proved to be permanent. The
20 activated electrode in FIGURE 1 was tested for about 4 weeks.
During this time, the cell was operated continuously for a few
days, shut down for a few days and then restarted the next
day, or frozen for a few days then started the next day.

Curves G and H in FIGURE 3 show the cell current densities at 0.70 and 0.50 V, respectively, when the cell was tested at 35/45/45°C. The performance fluctuated, but the trend showed very little decrease in 4 weeks.

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It is believed that the fluctuation was due mainly to water management rather than loss of activation. The cell was once frozen at -17°C for 3 days, after which it was difficult to start because the passes for the reactant flow were blocked by ice. After the ice was melted by the gaseous reactants, the cell immediately exhibited good performance.

Activation can be achieved very quickly. Referring to FIGURE 4, the cell performance during the activation process is shown at 75/95/90°C and 15/20 psig. The electrode had a Pt loading of 0.17 mg/cm² that was made using an old batch of Engelhard 20% Pt/C on a carbon paper-type gas diffusion medium. The current densities at 0.70 and 0.40 V, respectively, are shown in FIGURE 4 as curves I and J. The cell achieved 78% activation at 0.70 V, and 93% activation at 0.40 V, after only a short period (as little as 5 minutes). After 30 minutes, the cell achieved 87% activation at 0.70 V, and 97% activation at 0.40 V. After 60 minutes, the cell

achieved 93% activation at 0.70 V, and 100% activation at 0.40
V. After 90 minutes, the cell achieved 100% activation at
0.70 V. Clearly, the activation proceeded at a very fast pace
and most of the activation was achieved in the first few
5 minutes.

A variety of supported catalysts manufactured by
different companies was tested. They could all be fully
activated within 2 hours although different catalysts might
need different activation times. Generally speaking,
catalysts with lower Pt contents on the support (e.g., 10%
Pt/C) needed more activation time than those with higher Pt
contents on the support (e.g., 40% Pt/C). In addition, the
improvement in performance after activation is larger for
catalysts with lower Pt contents on the support. The
activation procedure was also applicable to electrodes made
using unsupported catalysts, such as Pt black, but the
increase in performance was normally less than that of
electrodes made using supported catalysts. The activation
proceeds faster at higher cell temperature and higher
20 backpressure. MEAs consisting of different kinds of membranes
or the same kind of membrane but having different thickness,
were all able to be activated quickly. The same is true for

MEAs whose electrodes contained various amounts of Nafion and PTFE.

5 The percentage performance improvement due to activation was different for electrodes with different Pt loadings, although they were made with the same catalyst. The lower Pt loading electrode shows more increase in performance.

10 Referring to FIGURE 5, the performances of six electrodes are shown. The electrodes were made using E-TEK 20% Pt/Vulcan XC-72 material with Pt loadings of 0.022, 0.043, 0.083, 0.137, 0.253, and 0.332 mg/cm², at 35/45/45°C before activation. The performance increased with Pt loading up to 0.253 mg/cm².

15 After being activated at 75/95/90°C and 20/30 psig for 2 hours, all of the electrodes showed large increases in performance, but the lower Pt loaded electrodes increased more in percentage, as shown in FIGURE 6. After activation, the performances of electrodes with Pt loadings of 0.083 to 0.332 mg/cm² were much closer with the electrode having a Pt loading of 0.138 mg/cm² exhibiting the best performance. It was found 20 that when 20% Pt/C was used to make an electrode, the Pt loading had to be controlled around 0.15 mg/cm² in order for

the best performance to be achieved with the minimum amount of Pt. Further increasing Pt loading increases the catalyst layer thickness, which in turn increases the electrical and ionic resistance of the catalyst layer and the mass transport 5 resistance to reactants.

Since other modifications and changes varied to fit particular operating requirements and environments will be apparent to those skilled in the art, the invention is not considered limited to the example chosen for purposes of disclosure, and covers all changes and modifications which do not constitute departures from the true spirit and scope of this invention.

Having thus described the invention, what is desired to be protected by Letters Patent is presented in the subsequently appended claims.

What is claimed is: